

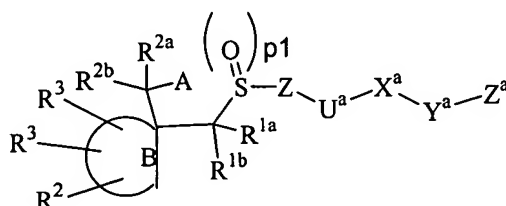
AMENDMENT

In the Claims:

Please cancel claims 7 and 15-18 without waiver or prejudice.

This listing of claims will replace all prior versions and listings of claims in the application.

1. (Previously presented) A compound of formula I:



I

or a stereoisomer or pharmaceutically acceptable salt form thereof, wherein;

A is selected from $-\text{COR}^5$, $-\text{CO}_2\text{H}$, $\text{CH}_2\text{CO}_2\text{H}$, $-\text{CO}_2\text{R}^6$, $-\text{CONHOH}$, $-\text{CONHOR}^5$, $-\text{CONHOR}^6$, $-\text{N}(\text{OH})\text{CHO}$, $-\text{N}(\text{OH})\text{COR}^5$, $-\text{SH}$, $-\text{CH}_2\text{SH}$, $-\text{SONHR}^a$, $-\text{SN}_2\text{H}_2\text{R}^a$, $-\text{PO}(\text{OH})_2$, and $-\text{PO}(\text{OH})\text{NHR}^a$;

ring B is a 5-6 membered heterocyclic ring consisting of: carbon atoms, 0-1 carbonyl groups, 0-1 double bonds, and from 0-2 ring heteroatoms selected from O, N, NR^2 , and $\text{S}(\text{O})_p$, provided that ring B contains other than a S-S, O-O, or S-O bond and provided that N- R^2 forms other than an N-O, N-N, or N-S bond;

Z is phenyl substituted with 0-4 R^b ;

U^a is O;

X^a is absent or selected from C_{1-10} alkylene, C_{2-10} alkenylene, and C_{2-10} alkynylene;

Y^a is absent or selected from O, NR^{a1} , $S(O)_p$, and $C(O)$;

Z^a is quinolinyl substituted with 0-5 R^c ;

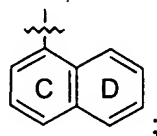
provided that Z, U^a , Y^a , and Z^a do not combine to form a O-N, O-O, or O- $S(O)_p$ group;

R^{1a} is selected from H, C_{1-4} alkyl, phenyl, benzyl, CH_2OR^3 , and $CH_2NR^aR^{a1}$;

R^{1b} is selected from H, C_{1-4} alkyl, phenyl, benzyl, CH_2OR^3 , and $CH_2NR^aR^{a1}$;

alternatively, R^{1a} and R^{1b} combine to form a 3-6 membered ring consisting of: carbon atoms and 0-1 heteroatoms selected from O, S, $S(O)$, $S(O)_2$, and NR^a ;

provided that when R^{1a} and R^{1b} are hydrogen and ring B is a heterocycle, then Z^a is the following:



ring C is phenyl or pyridyl and is substituted with 0-2 R^c ;

ring D is selected from phenyl, pyridyl, pyridazinyl, pyrimidyl, and pyrazinyl, and is substituted with 0-3 R^c ;

R^2 is selected from Q, C_{1-10} alkylene-Q substituted with 0-3 R^{b1} , C_{2-10} alkenylene-Q substituted with 0-3 R^{b1} , C_{2-10} alkynylene-Q substituted with 0-3 R^{b1} , $(CR^aR^{a1})_{r1}O(CR^aR^{a1})_{r-Q}$, $(CR^aR^{a1})_{r1}NR^a(CR^aR^{a1})_{r-Q}$, $(CR^aR^{a1})_{r1}C(O)(CR^aR^{a1})_{r-Q}$, $(CR^aR^{a1})_{r1}C(O)O(CR^aR^{a1})_{r-Q}$,

$(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{OC}(\text{O})(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}}$, $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{C}(\text{O})\text{NR}^{\text{aR}^{\text{a1}}}$,
 $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{C}(\text{O})\text{NR}^{\text{a}}(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}}$, $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{NR}^{\text{a}}\text{C}(\text{O})(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}}$,
 $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{OC}(\text{O})\text{O}(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}}$, $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{OC}(\text{O})\text{NR}^{\text{a}}(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}}$,
 $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{NR}^{\text{a}}\text{C}(\text{O})\text{O}(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}}$, $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{NR}^{\text{a}}\text{C}(\text{O})\text{NR}^{\text{a}}(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}}$,
 $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{S}(\text{O})_{\text{p}}(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}}$, $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{SO}_2\text{NR}^{\text{a}}(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}}$,
 $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{NR}^{\text{a}}\text{SO}_2(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}}$, and $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{NR}^{\text{a}}\text{SO}_2\text{NR}^{\text{a}}(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}}$;

$\text{R}^{2\text{a}}$ is selected from H, C_{1-4} alkyl, phenyl, benzyl, CH_2OR^3 , and $\text{CH}_2\text{NR}^{\text{aR}^{\text{a1}}}$;

$\text{R}^{2\text{b}}$ is selected from H, C_{1-4} alkyl, phenyl, benzyl, CH_2OR^3 , and $\text{CH}_2\text{NR}^{\text{aR}^{\text{a1}}}$;

alternatively, $\text{R}^{2\text{a}}$ and $\text{R}^{2\text{b}}$ combine to form a 3-6 membered ring consisting of: carbon atoms and 0-1 heteroatoms selected from O, S, $\text{S}(\text{O})$, $\text{S}(\text{O})_2$, and NR^{a} ;

Q is selected from H, a C_{3-13} carbocyclic residue substituted with 0-5 R^{d} and a 5-14 membered heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $\text{S}(\text{O})_{\text{p}}$ and substituted with 0-5 R^{d} ;

R^3 , at each occurrence, is selected from Q^1 , C_{1-6} alkylene- Q^1 , C_{2-6} alkenylene- Q^1 , C_{2-6} alkynylene- Q^1 , $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{O}(\text{CH}_2)_{\text{r-Q}^1}$, $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{NR}^{\text{a}}(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}^1}$,
 $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{NR}^{\text{a}}\text{C}(\text{O})(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}^1}$, $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{C}(\text{O})\text{NR}^{\text{a}}(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}^1}$,
 $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{C}(\text{O})(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}^1}$, $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{C}(\text{O})\text{O}(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}^1}$,
 $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{S}(\text{O})_{\text{p}}(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}^1}$, and $(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r1}}\text{SO}_2\text{NR}^{\text{a}}(\text{CR}^{\text{aR}^{\text{a1}}})_{\text{r-Q}^1}$;

alternatively, when two R^3 's are attached to the same carbon atom, they combine to form a 3-8 membered carbocyclic or heterocyclic ring consisting of: carbon atoms and

0-3 heteroatoms selected from the group consisting of N, O, and S(O)_p and substituted with 0-3 R^d;

Q¹ is selected from H, phenyl substituted with 0-3 R^d, naphthyl substituted with 0-3 R^d and a 5-10 membered heteroaryl consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S and substituted with 0-3 R^d;

R^a, at each occurrence, is independently selected from H, C₁₋₄ alkyl, phenyl and benzyl;

R^{a1}, at each occurrence, is independently selected from H and C₁₋₄ alkyl;

alternatively, R^a and R^{a1} when attached to a nitrogen are taken together with the nitrogen to which they are attached to form a 5 or 6 membered ring comprising carbon atoms and from 0-1 additional heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{a2}, at each occurrence, is independently selected from C₁₋₄ alkyl, phenyl and benzyl;

R^b, at each occurrence, is independently selected from C₁₋₆ alkyl, OR^a, Cl, F, Br, I, =O, -CN, NO₂, NR^aR^{a1}, C(O)R^a, C(O)OR^a, C(O)NR^aR^{a1}, R^aNC(O)NR^aR^{a1}, OC(O)NR^aR^{a1}, R^aNC(O)O, S(O)₂NR^aR^{a1}, NR^aS(O)₂R^{a2}, NR^aS(O)₂NR^aR^{a1}, OS(O)₂NR^aR^{a1}, NR^aS(O)₂R^{a2}, S(O)_pR^{a2}, CF₃, and CF₂CF₃;

R^{b1}, at each occurrence, is independently selected from OR^a, Cl, F, Br, I, =O, -CN, NO₂, and NR^aR^{a1};

R^c, at each occurrence, is independently selected from C₁₋₆ alkyl, OR^a, Cl, F, Br, I, =O, -CN, NO₂, NR^aR^{a1}, C(O)R^a, C(O)OR^a, C(O)NR^aR^{a1}, R^aNC(O)NR^aR^{a1},

OC(O)NR^aR^{a1}, R^aNC(O)O, S(O)₂NR^aR^{a1}, NR^aS(O)₂R^{a2}, NR^aS(O)₂NR^aR^{a1}, OS(O)₂NR^aR^{a1}, NR^aS(O)₂R^{a2}, S(O)_pR^{a2}, CF₃, CF₂CF₃, C₃₋₁₀ carbocyclic residue and a 5-14 membered heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^d, at each occurrence, is independently selected from C₁₋₆ alkyl, OR^a, Cl, F, Br, I, =O, -CN, NO₂, NR^aR^{a1}, C(O)R^a, C(O)OR^a, C(O)NR^aR^{a1}, R^aNC(O)NR^aR^{a1}, OC(O)NR^aR^{a1}, R^aNC(O)O, S(O)₂NR^aR^{a1}, NR^aS(O)₂R^{a2}, NR^aS(O)₂NR^aR^{a1}, OS(O)₂NR^aR^{a1}, NR^aS(O)₂R^{a2}, S(O)_pR^{a2}, CF₃, CF₂CF₃, C₃₋₁₀ carbocyclic residue and a 5-14 membered heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R⁵, at each occurrence, is selected from C₁₋₁₀ alkyl substituted with 0-2 R^b, and C₁₋₈ alkyl substituted with 0-2 R^e;

R^e, at each occurrence, is selected from phenyl substituted with 0-2 R^b and biphenyl substituted with 0-2 R^b;

R⁶, at each occurrence, is selected from phenyl, naphthyl, C₁₋₁₀ alkyl-phenyl-C₁₋₆ alkyl-, C₃₋₁₁ cycloalkyl, C₁₋₆ alkylcarbonyloxy-C₁₋₃ alkyl-, C₁₋₆ alkoxy carbonyloxy-C₁₋₃ alkyl-, C₂₋₁₀ alkoxy carbonyl, C₃₋₆ cycloalkylcarbonyloxy-C₁₋₃ alkyl-, C₃₋₆ cycloalkoxy carbonyloxy-C₁₋₃ alkyl-, C₃₋₆ cycloalkoxy carbonyl, phenoxycarbonyl, phenyloxy carbonyloxy-C₁₋₃ alkyl-, phenylcarbonyloxy-C₁₋₃ alkyl-, C₁₋₆ alkoxy-C₁₋₆ alkylcarbonyloxy-C₁₋₃ alkyl-, [5-(C₁-C₅ alkyl)-1,3-dioxo-cyclopenten-2-one-yl]methyl, [5-(R^a)-1,3-dioxo-cyclopenten-2-one-yl]methyl, (5-aryl-1,3-dioxo-cyclopenten-2-one-yl)methyl, -C₁₋₁₀ alkyl-NR⁷R^{7a}, -CH(R⁸)OC(=O)R⁹, and -CH(R⁸)OC(=O)OR⁹;

R⁷ is selected from H and C₁₋₁₀ alkyl, C₂₋₆ alkenyl, C₃₋₆ cycloalkyl-C₁₋₃ alkyl-, and phenyl-C₁₋₆ alkyl-;

R^{7a} is selected from H and C₁₋₁₀ alkyl, C₂₋₆ alkenyl, C₃₋₆ cycloalkyl-C₁₋₃ alkyl-, and phenyl-C₁₋₆ alkyl-;

R⁸ is selected from H and C₁₋₄ linear alkyl;

R⁹ is selected from H, C₁₋₈ alkyl substituted with 1-2 R^f, C₃₋₈ cycloalkyl substituted with 1-2 R^f, and phenyl substituted with 0-2 R^b;

R^f, at each occurrence, is selected from C₁₋₄ alkyl, C₃₋₈ cycloalkyl, C₁₋₅ alkoxy, and phenyl substituted with 0-2 R^b;

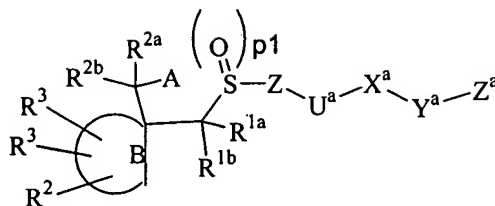
p, at each occurrence, is selected from 0, 1, and 2;

p₁ is selected from 0, 1, and 2;

r, at each occurrence, is selected from 0, 1, 2, 3, and 4; and,

r₁, at each occurrence, is selected from 0, 1, 2, 3, and 4.

2. (Previously presented) A compound according to Claim 1, wherein the compound is of formula II:



II

or a stereoisomer or pharmaceutically acceptable salt form thereof, wherein;

A is selected from $-\text{CO}_2\text{H}$, $\text{CH}_2\text{CO}_2\text{H}$, $-\text{CONHOH}$, $-\text{CONHOR}^5$, $-\text{CONHOR}^6$, $-\text{N}(\text{OH})\text{CHO}$, $-\text{N}(\text{OH})\text{COR}^5$, $-\text{SH}$, and $-\text{CH}_2\text{SH}$;

ring B is a 5-6 membered heterocyclic ring consisting of: carbon atoms, 0-1 carbonyl groups, 0-1 double bonds, and from 0-2 ring heteroatoms selected from O, N, and NR^2 , provided that ring B contains other than an O-O bond and provided that NR^2 forms other than an N-O, N-N, or N-S bond;

X^a is absent or selected from C_{1-4} alkylene and C_{2-4} alkynylene;

Y^a is absent or selected from O and NR^{a1} ;

provided that Z, U^a , Y^a , and Z^a do not combine to form a O-N or O-O group;

R^2 is selected from Q, C_{1-6} alkylene-Q, C_{2-6} alkenylene-Q, C_{2-6} alkynylene-Q, $(\text{CR}^a\text{R}^{a1})_{r1}\text{O}(\text{CR}^a\text{R}^{a1})_{r-Q}$, $(\text{CR}^a\text{R}^{a1})_{r1}\text{NR}^a(\text{CR}^a\text{R}^{a1})_{r-Q}$, $(\text{CR}^a\text{R}^{a1})_{r1}\text{C}(\text{O})(\text{CR}^a\text{R}^{a1})_{r-Q}$, $(\text{CR}^a\text{R}^{a1})_{r1}\text{C}(\text{O})\text{O}(\text{CR}^a\text{R}^{a1})_{r-Q}$, $(\text{CR}^a\text{R}^{a1})_{r1}\text{C}(\text{O})\text{NR}^a\text{R}^{a1}$, $(\text{CR}^a\text{R}^{a1})_{r1}\text{C}(\text{O})\text{NR}^a(\text{CR}^a\text{R}^{a1})_{r-Q}$, $(\text{CR}^a\text{R}^{a1})_{r1}\text{S}(\text{O})_p(\text{CR}^a\text{R}^{a1})_{r-Q}$, and $(\text{CR}^a\text{R}^{a1})_{r1}\text{SO}_2\text{NR}^a(\text{CR}^a\text{R}^{a1})_{r-Q}$;

Q is selected from H, a C₃₋₆ carbocyclic residue substituted with 0-5 R^d, and a 5-10 membered heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p and substituted with 0-5 R^d;

R^a, at each occurrence, is independently selected from H, C₁₋₄ alkyl, phenyl and benzyl;

R^{a1}, at each occurrence, is independently selected from H and C₁₋₄ alkyl;

alternatively, R^a and R^{a1} when attached to a nitrogen are taken together with the nitrogen to which they are attached to form a 5 or 6 membered ring comprising carbon atoms and from 0-1 additional heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^{a2}, at each occurrence, is independently selected from C₁₋₄ alkyl, phenyl and benzyl;

R^b, at each occurrence, is independently selected from C₁₋₆ alkyl, OR^a, Cl, F, Br, =O, -CN, NR^aR^{a1}, C(O)R^a, C(O)OR^a, C(O)NR^aR^{a1}, S(O)₂NR^aR^{a1}, S(O)_pR^{a2}, and CF₃;

R^c, at each occurrence, is independently selected from C₁₋₆ alkyl, OR^a, Cl, F, Br, =O, -CN, NR^aR^{a1}, C(O)R^a, C(O)OR^a, C(O)NR^aR^{a1}, S(O)₂NR^aR^{a1}, S(O)_pR^{a2}, CF₃, C₃₋₆ carbocyclic residue and a 5-6 membered heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R^d, at each occurrence, is independently selected from C₁₋₆ alkyl, OR^a, Cl, F, Br, =O, -CN, NR^aR^{a1}, C(O)R^a, C(O)OR^a, C(O)NR^aR^{a1}, S(O)₂NR^aR^{a1}, S(O)_pR^{a2}, CF₃, C₃₋₆ carbocyclic residue and a 5-6 membered heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and S(O)_p;

R⁵, at each occurrence, is selected from C₁₋₆ alkyl substituted with 0-2 R^b, and C₁₋₄ alkyl substituted with 0-2 R^e;

R^e, at each occurrence, is selected from phenyl substituted with 0-2 R^b and biphenyl substituted with 0-2 R^b;

R⁶, at each occurrence, is selected from phenyl, naphthyl, C₁₋₁₀ alkyl-phenyl-C₁₋₆ alkyl-, C₃₋₁₁ cycloalkyl, C₁₋₆ alkylcarbonyloxy-C₁₋₃ alkyl-, C₁₋₆ alkoxycarbonyloxy-C₁₋₃ alkyl-, C₂₋₁₀ alkoxycarbonyl, C₃₋₆ cycloalkylcarbonyloxy-C₁₋₃ alkyl-, C₃₋₆ cycloalkoxycarbonyloxy-C₁₋₃ alkyl-, C₃₋₆ cycloalkoxycarbonyl, phenoxycarbonyl, phenyloxycarbonyloxy-C₁₋₃ alkyl-, phenylcarbonyloxy-C₁₋₃ alkyl-, C₁₋₆ alkoxy-C₁₋₆ alkylcarbonyloxy-C₁₋₃ alkyl-, [5-(C₁-C₅ alkyl)-1,3-dioxo-cyclopenten-2-one-yl]methyl, [5-(R^a)-1,3-dioxo-cyclopenten-2-one-yl]methyl, (5-aryl-1,3-dioxo-cyclopenten-2-one-yl)methyl, -C₁₋₁₀ alkyl-NR⁷R^{7a}, -CH(R⁸)OC(=O)R⁹, and -CH(R⁸)OC(=O)OR⁹;

R⁷ is selected from H and C₁₋₆ alkyl, C₂₋₆ alkenyl, C₃₋₆ cycloalkyl-C₁₋₃ alkyl-, and phenyl-C₁₋₆ alkyl-;

R^{7a} is selected from H and C₁₋₆ alkyl, C₂₋₆ alkenyl, C₃₋₆ cycloalkyl-C₁₋₃ alkyl-, and phenyl-C₁₋₆ alkyl-;

R⁸ is selected from H and C₁₋₄ linear alkyl;

R⁹ is selected from H, C₁₋₆ alkyl substituted with 1-2 R^f, C₃₋₆ cycloalkyl substituted with 1-2 R^f, and phenyl substituted with 0-2 R^b;

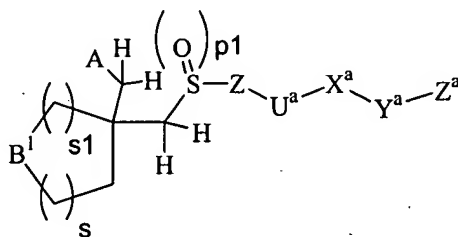
R^f , at each occurrence, is selected from C_{1-4} alkyl, C_{3-6} cycloalkyl, C_{1-5} alkoxy, and phenyl substituted with 0-2 R^b ;

p , at each occurrence, is selected from 0, 1, and 2;

r , at each occurrence, is selected from 0, 1, 2, 3, and 4; and,

r_1 , at each occurrence, is selected from 0, 1, 2, 3, and 4.

3. (Previously presented) A compound according to Claim 2, wherein the compound is of formula III:



III

or a stereoisomer or pharmaceutically acceptable salt form thereof, wherein;

A is selected from $-CO_2H$, CH_2CO_2H , $-CONHOH$, $-CONHOR^5$, $-N(OH)CHO$, and $-N(OH)COR^5$;

B^1 is NR^2 or O;

Z is phenyl substituted with 0-3 R^b ;

X^a is absent or selected from C_{1-2} alkylene and C_{2-4} alkynylene;

R^2 is selected from Q, C_{1-6} alkylene-Q, C_{2-6} alkenylene-Q, C_{2-6} alkynylene-Q,

$(CR^aR^{a1})_{r1}O(CR^aR^{a1})_{r-Q}$, $(CR^aR^{a1})_{r1}NR^a(CR^aR^{a1})_{r-Q}$,

$(CR^aR^{a1})_{r1}C(O)(CR^aR^{a1})_{r-Q}$, $(CR^aR^{a1})_{r1}C(O)O(CR^aR^{a1})_{r-Q}$,

$(CR^aR^{a2})_{r1}C(O)NR^aR^{a1}$, $(CR^aR^{a2})_{r1}C(O)NR^a(CR^aR^{a1})_{r-Q}$, and

$(CR^aR^{a1})_{r1}S(O)_p(CR^aR^{a1})_{r-Q}$;

Q is selected from H, a C_{3-6} carbocyclic residue substituted with 0-3 R^d and a 5-10 membered heterocycle consisting of: carbon atoms and 1-4 heteroatoms selected from the group consisting of N, O, and $S(O)_p$ and substituted with 0-3 R^d ;

R^a , at each occurrence, is independently selected from H, C_{1-4} alkyl, phenyl and benzyl;

R^{a1} , at each occurrence, is independently selected from H and C_{1-4} alkyl;

R^{a2} , at each occurrence, is independently selected from C_{1-4} alkyl, phenyl and benzyl;

R^b , at each occurrence, is independently selected from C_{1-4} alkyl, OR^a , Cl, F, =O, NR^aR^{a1} , $C(O)R^a$, $C(O)OR^a$, $C(O)NR^aR^{a1}$, $S(O)_2NR^aR^{a1}$, $S(O)_pR^{a2}$, and CF_3 ;

R^c , at each occurrence, is independently selected from C_{1-6} alkyl, OR^a , Cl, F, Br, =O, NR^aR^{a1} , $C(O)R^a$, $C(O)NR^aR^{a1}$, $S(O)_2NR^aR^{a1}$, $S(O)_pR^{a2}$, and CF_3 ;

R^d , at each occurrence, is independently selected from C_{1-6} alkyl, OR^a , Cl, F, Br, =O, NR^aR^{a1} , $C(O)R^a$, $C(O)NR^aR^{a1}$, $S(O)_2NR^aR^{a1}$, $S(O)_pR^{a2}$, CF_3 and phenyl;

R^5 , at each occurrence, is selected from C_{1-4} alkyl substituted with 0-2 R^b , and C_{1-4} alkyl substituted with 0-2 R^c ;

R^e , at each occurrence, is selected from phenyl substituted with 0-2 R^b and biphenyl substituted with 0-2 R^b ;

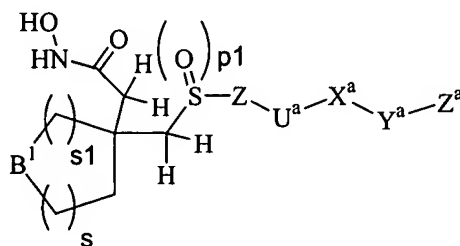
p , at each occurrence, is selected from 0, 1, and 2;

r , at each occurrence, is selected from 0, 1, 2, 3, and 4;

$r1$, at each occurrence, is selected from 0, 1, 2, 3, and 4; and,

s and $s1$ combine to total 1, 2, 3, or 4.

4. (Previously presented) A compound according to Claim 3, wherein the compound is of formula IV:



IV

or a stereoisomer or pharmaceutically acceptable salt form thereof, wherein;

Z is phenyl substituted with 0-3 R^b ;

X^a is absent or is selected from CH_2 , CH_2CH_2 , and C_{2-4} alkynylene;

Y^a is absent or is O;

Z^a is quinolinyl substituted with 0-3 R^c ;

provided that Z, U^a, Y^a, and Z^a do not combine to form a O-O group;

R² is selected from Q, C₁₋₆ alkylene-Q, C₂₋₆ alkynylene-Q, (CR^aR^{a1})_{r1}O(CR^aR^{a1})_{r-Q}, (CR^aR^{a1})_{r1}NR^a(CR^aR^{a1})_{r-Q}, C(O)(CR^aR^{a1})_{r-Q}, C(O)O(CR^aR^{a1})_{r-Q}, C(O)NR^a(CR^aR^{a1})_{r-Q}, and S(O)_p(CR^aR^{a1})_{r-Q};

Q is selected from H, cyclopropyl substituted with 0-1 R^d, cyclobutyl substituted with 0-1 R^d, cyclopentyl substituted with 0-1 R^d, cyclohexyl substituted with 0-1 R^d, phenyl substituted with 0-2 R^d and a heteroaryl substituted with 0-3 R^d, wherein the heteroaryl is selected from pyridyl, quinolinyl, thiazolyl, furanyl, imidazolyl, and isoxazolyl;

R^a, at each occurrence, is independently selected from H, CH₃, and CH₂CH₃;

R^{a1}, at each occurrence, is independently selected from H, CH₃, and CH₂CH₃;

R^{a2}, at each occurrence, is independently selected from H, CH₃, and CH₂CH₃;

R^b, at each occurrence, is independently selected from C₁₋₄ alkyl, OR^a, Cl, F, =O, NR^aR^{a1}, C(O)R^a, C(O)OR^a, C(O)NR^aR^{a1}, S(O)₂NR^aR^{a1}, S(O)_pR^{a2}, and CF₃;

R^c, at each occurrence, is independently selected from C₁₋₆ alkyl, OR^a, Cl, F, Br, =O, NR^aR^{a1}, C(O)R^a, C(O)NR^aR^{a1}, S(O)₂NR^aR^{a1}, S(O)_pR^{a2}, and CF₃;

R^d, at each occurrence, is independently selected from C₁₋₆ alkyl, OR^a, Cl, F, Br, =O, NR^aR^{a1}, C(O)R^a, C(O)NR^aR^{a1}, S(O)₂NR^aR^{a1}, S(O)_pR^{a2}, CF₃ and phenyl;

p, at each occurrence, is selected from 0, 1, and 2;

r, at each occurrence, is selected from 0, 1, 2, and 3;

r1, at each occurrence, is selected from 0, 1, 2, and 3; and,

s and s1 combine to total 2, 3, or 4.

5. (Previously presented) A compound according to Claim 1, wherein the compound is selected from the group:

N-hydroxy-2-{2-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-2-pyrrolidinyl}acetamide;

N-hydroxy-2-{1-methyl-2-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-2-pyrrolidinyl}acetamide;

N-hydroxy-2-{1-isobutyl-2-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-2-pyrrolidinyl}acetamide;

N-hydroxy-2-[2-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-1-(3-pyridinyl)-2-pyrrolidinyl}acetamide;

2-{1-acetyl-2-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-2-pyrrolidinyl}-*N*-hydroxyacetamide;

N-hydroxy-2-{3-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-3-pyrrolidinyl}acetamide;

N-hydroxy-2-{1-methyl-3-[(4-{(2-methyl-4-quinolinyl)methoxy}phenyl)sulfonyl)methyl]-3-pyrrolidinyl}acetamide;

N-hydroxy-2-{1-isopropyl-3-[(4-{(2-methyl-4-quinolinyl)methoxy}phenyl)sulfonyl)methyl]-3-pyrrolidinyl}acetamide;

N-hydroxy-2-{1-isobutyl-3-[(4-{(2-methyl-4-quinolinyl)methoxy}phenyl)sulfonyl)methyl]-3-pyrrolidinyl}acetamide;

N-hydroxy-2-{3-[(4-{(2-methyl-4-quinolinyl)methoxy}phenyl)sulfonyl)methyl]-1-neopentyl-3-pyrrolidinyl}acetamide;

N-hydroxy-2-{2-[(4-{(2-methyl-4-quinolinyl)methoxy}phenyl)sulfonyl)methyl]-2-piperidinyl}acetamide;

N-hydroxy-2-{1-methyl-2-[(4-{(2-methyl-4-quinolinyl)methoxy}phenyl)sulfonyl)methyl]-2-piperidinyl}acetamide;

N-hydroxy-2-{1-isobutyl-2-[(4-{(2-methyl-4-quinolinyl)methoxy}phenyl)sulfonyl)methyl]-2-piperidinyl}acetamide;

N-hydroxy-2-{3-[(4-{(2-methyl-4-quinolinyl)methoxy}phenyl)sulfonyl)methyl]-3-piperidinyl}acetamide;

N-hydroxy-2-{1-methyl-3-[(4-{(2-methyl-4-quinolinyl)methoxy}phenyl)sulfonyl)methyl]-3-piperidinyl}acetamide;

N-hydroxy-2-{1-isopropyl-3-[(4-{(2-methyl-4-quinolinyl)methoxy}phenyl)sulfonyl)methyl]-3-piperidinyl}acetamide;

N-hydroxy-2-{3-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-3-piperidinyl}acetamide;

N-hydroxy-2-{1-methyl-3-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-3-piperidinyl}acetamide;

N-hydroxy-2-{1-isopropyl-3-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-3-piperidinyl}acetamide;

N-hydroxy-2-{1-isobutyl-3-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-3-piperidinyl}acetamide;

N-hydroxy-2-{4-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-4-piperidinyl}acetamide;

N-hydroxy-2-{1-methyl-4-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-4-piperidinyl}acetamide;

N-hydroxy-2-{2-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]tetrahydro-2-furanyl}acetamide;

N-hydroxy-2-{1-methyl-3-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-2-oxopyrrolidinyl}acetamide;

N-hydroxy-2-[5-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-3-(3-pyridinyl)-4,5-dihydro-5-isoxazolyl]acetamide;

N-hydroxy-2-[5-[(4-[(2-methyl-4-quinolinyl)methoxy]phenyl)sulfonyl)methyl]-3-(4-pyridinyl)-4,5-dihydro-5-isoxazolyl]acetamide; and,

N-hydroxy-2-{4-[(4-{(2-methyl-4-quinolinyl)methoxy}phenyl)sulfonyl)methyl]tetrahydro-2*H*-pyran-4-yl}acetamide;

or a pharmaceutically acceptable salt form thereof.

6. (Original) A pharmaceutical composition, comprising: a pharmaceutically acceptable carrier and a therapeutically effective amount of a compound according to Claim 1 or a pharmaceutically acceptable salt form thereof.

7-10. (Canceled).

11. (Previously presented) A pharmaceutical composition, comprising: a pharmaceutically acceptable carrier and a therapeutically effective amount of a compound according to Claim 2 or a pharmaceutically acceptable salt form thereof.

12. (Previously presented) A pharmaceutical composition, comprising: a pharmaceutically acceptable carrier and a therapeutically effective amount of a compound according to Claim 3 or a pharmaceutically acceptable salt form thereof.

13. (Previously presented) A pharmaceutical composition, comprising: a pharmaceutically acceptable carrier and a therapeutically effective amount of a compound according to Claim 4 or a pharmaceutically acceptable salt form thereof.

14. (Previously presented) A pharmaceutical composition, comprising: a pharmaceutically acceptable carrier and a therapeutically effective amount of a compound according to Claim 5 or a pharmaceutically acceptable salt form thereof.

15-26. (Canceled).

27. (Previously presented) A method of treating a disease or condition in a patient, comprising: administering to the mammal in need of such treatment a therapeutically effective amount of a compound of Claim 1 or a pharmaceutically acceptable salt form thereof, wherein the disease or condition is selected from acute infection, acute phase response, allergic asthma, anorexia, asthma, autoimmune disease, cachexia, cardiovascular effects, coagulation, fever, gingivitis, graft versus host disease, hemorrhage, multiple sclerosis, neovascular glaucoma, osteoarthritis, periodontitis, psoriasis, psoriatic arthritis, rheumatic fever, rheumatoid arthritis, shock, and solid tumor growth and tumor invasion by secondary metastases.

28. (Previously presented) A method of treating a disease or condition in a patient, comprising: administering to the mammal in need of such treatment a therapeutically effective amount of a compound of Claim 2 or a pharmaceutically acceptable salt form thereof, wherein the disease or condition is selected from acute infection, acute phase response, allergic asthma, anorexia, asthma, autoimmune disease, cachexia, cardiovascular effects, coagulation, fever, gingivitis, graft versus host disease, hemorrhage, multiple sclerosis, neovascular glaucoma, osteoarthritis, periodontitis, psoriasis, psoriatic arthritis, rheumatic fever, rheumatoid arthritis, shock, and solid tumor growth and tumor invasion by secondary metastases.

29. (Previously presented) A method of treating a disease or condition in a patient, comprising: administering to the mammal in need of such treatment a therapeutically effective amount of a compound of Claim 3 or a pharmaceutically acceptable salt form thereof, wherein the disease or condition is selected from acute infection, acute phase response, allergic asthma, anorexia, asthma, autoimmune disease, cachexia, cardiovascular effects, coagulation, fever, gingivitis, graft versus host disease, hemorrhage, multiple sclerosis, neovascular glaucoma, osteoarthritis, periodontitis, psoriasis, psoriatic arthritis, rheumatic fever, rheumatoid arthritis, shock, and solid tumor growth and tumor invasion by secondary metastases.

30. (Previously presented) A method of treating a disease or condition in a patient, comprising: administering to the mammal in need of such treatment a therapeutically effective amount of a compound of Claim 4 or a pharmaceutically acceptable salt form thereof, wherein the disease or condition is selected from acute infection, acute phase response, allergic asthma, anorexia, asthma, autoimmune disease, cachexia, cardiovascular effects, coagulation, fever, gingivitis, graft versus host disease, hemorrhage, multiple sclerosis, neovascular glaucoma, osteoarthritis, periodontitis, psoriasis, psoriatic arthritis, rheumatic fever, rheumatoid arthritis, shock, and solid tumor growth and tumor invasion by secondary metastases.

31. (Previously presented) A method of treating a disease or condition in a patient, comprising: administering to the mammal in need of such treatment a therapeutically effective amount of a compound of Claim 5 or a pharmaceutically acceptable salt form thereof, wherein the disease or condition is selected from acute infection, acute phase response, allergic asthma, anorexia, asthma, autoimmune disease, cachexia, cardiovascular effects, coagulation, fever, gingivitis, graft versus host disease, hemorrhage, multiple sclerosis, neovascular glaucoma, osteoarthritis, periodontitis, psoriasis, psoriatic arthritis, rheumatic fever, rheumatoid arthritis, shock, and solid tumor growth and tumor invasion by secondary metastases.